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TAF SPRAY 28.5 MG/G CUTANEOUS SPRAY, SOLUTION

Ima dovoljenje za promet

- Thiamphenicol

Informacije o zdravilu

Ime zdravila:
TAF SPRAY 28.5 MG/G CUTANEOUS SPRAY, SOLUTION

Učinkovina:
Na voljo samo v English

Ciljne živalske vrste:
govedo
Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian
Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian
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Pot uporabe:
Dermalna uporaba

Podatki o zdravilu

Učinkovina / Jakost:
Na voljo samo v English
28.50 milligram(s) / 1.00 gram(s)

Farmacevtska oblika:
Dermalno pršilo, raztopina

Karenca glede na pot uporabe:
Dermalna uporaba:

- **govedo**
  - Milk. 0 day

Do not use on the udder of lactating animals if their milk is intended for human consumption.

  - Meat and offal. 0 day

- **Pig**
  - Meat and offal. 14 day

- **Rabbit**
  - Meat and offal. 0 day

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Equid

- Meat and offal. 0 day

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Sheep

- Meat and offal. 0 day

- Milk. 0 day

Do not use on the udder of lactating animals if their milk is intended for human consumption.

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Goat

- Milk. 0 day

Do not use on the udder of lactating animals if their milk is intended for human consumption.

- Meat and offal. 0 day

Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):
QD06AX

Pravni status za dobavo/izdajo zdravila:
Zdravilo, ki se izdaja na veterinarski recept

Status dovoljenja za promet z zdravilom:
Valid

Ima dovoljenje za promet v:
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

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Na voljo v:
Belgium

Opis ovojnine zdravila:
Na voljo samo v French
Na voljo samo v French

Na voljo samo v French
Na voljo samo v French
Na voljo samo v French

Dodatne informacije

Vrsta dovoljenja:
Na voljo samo v English French Croatian Italian Latvian Finnish Swedish Icelandic Norwegian

Pravna podlaga za izdajo dovoljenja za promet z zdravilom:
Na voljo samo v English Italian Latvian Norwegian

Imetnik dovoljenja za promet z zdravilom:
Dechra Regulatory B.V.

Datum dovoljenja za promet z zdravilom:
6/01/2015

Proizvodna mesta, odgovorna za sproščanje serij:
Eurovet Animal Health B.V.
IGS Aerosols GmbH

Pristojni organ:
Federal Agency For Medicines And Health Products

Številka dovoljenja za promet z zdravilom:
BE-V467937

Datum spremembe statusa dovoljenja za promet:
6/01/2015

Referenčna država članica:
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

Številka postopka:
FR/V/0276/001

Zadevne države članice:
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
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Na voljo samo v Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

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Za poročila o domnevnih neželenih učinkih zdravil za uporabo v veterinaski medicini obiščite stran: www.adrreports.eu/vet

Dokumenti

Povzetek glavnih značilnosti zdravila

Ta dokument ne obstaja v tem jeziku (slovenščina). Lahko ga najdete spodaj v drugem jeziku.

Navodilo za uporabo

Ta dokument ne obstaja v tem jeziku (slovenščina). Lahko ga najdete spodaj v drugem jeziku.

Označevanje

Ta dokument ne obstaja v tem jeziku (slovenščina). Lahko ga najdete spodaj v drugem jeziku.

Combined File of all Documents

Ta dokument ne obstaja v tem jeziku (slovenščina). Lahko ga najdete spodaj v drugem jeziku.

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